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## **AMENDMENTS TO THE CLAIMS**

1. (Original) A triazolopyrimidine of the formula I

$$N - N$$
 $R^1$ 
 $R^2$ 

in which the substituents are as defined below:

$$R^1$$
 is  $C_1$ - $C_5$ -alkyl or  $C_1$ - $C_{10}$ -alkoxy- $C_1$ - $C_{10}$ -alkyl,

$$R^2$$
 is  $C_5$ - $C_{12}$ -alkyl,

where R<sup>1</sup> and/or R<sup>2</sup> may be substituted by one to three of the following groups:

cyano, nitro, hydroxyl, C<sub>3</sub>-C<sub>6</sub>-cycloalkyl, C<sub>1</sub>-C<sub>6</sub>-alkylthio, NR<sup>a</sup>R<sup>b</sup>;

 $R^a$ ,  $R^b$  are hydrogen or  $C_1$ - $C_6$ -alkyl.

- 2. (Original) The compound of the formula I according to claim 1 in which R<sup>1</sup> and R<sup>2</sup> are unsubstituted and together have at most 14 carbon atoms.
- 3. (Original) The compound of the formula I according to claim 1 in which R<sup>1</sup> is methyl, ethyl,

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n-propyl, isopropyl, n-butyl or n-pentyl, where the carbon chains are unsubstituted or may be substituted according to claim 1.

- 4. (Original) The compound of the formula I according to claim 1 in which R<sup>2</sup> is n-heptyl, n-octyl, n-nonyl or 1-methyloctyl.
- 5. (Original) 6-Methyl-5-pentyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;
  - 5-Hexyl-6-methyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;
  - 5-Heptyl-6-methyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;
  - 6-Methyl-5-octyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;
  - 6-Methyl-5-nonyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;
  - 6-Ethyl-5-octyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;
  - 6-Ethyl-5-noctyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;
    - 5-Decyl-6-ethyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;
    - 5-Octyl-6-propyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;
    - 5-Nonyl-6-propyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;
    - 5-Decyl-6-propyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine.
- 6. (Currently amended) A process for preparing compounds of the formula I according to any of claims 1 to 5 claim 1 wherein β-keto esters of the formula II,

$$RO$$
 $R^1$ 
 $R^2$ 
 $R^2$ 
 $R^3$ 

in which R is C<sub>1</sub>-C<sub>4</sub>-alkyl are reacted with 3-amino-1,2,4-triazole of the formula III

to give 7-hydroxytriazolopyrimidines of the formula IV

$$\begin{array}{c|c}
 & OH \\
 & N \\
 & N \\
 & N \\
 & R^2
\end{array}$$
IV

which are halogenated to give compounds of the formula V

$$\begin{array}{c|c}
 & \text{Hal} \\
 & \text{N} \\
 & \text{N} \\
 & \text{N} \\
 & \text{R}^2
\end{array}$$

in which Hal is chlorine or bromine, and V is reacted with ammonia.

7. (Currently amended) A process for preparing compounds of the formula I according to any of claims 1 to 5 claim 1 wherein acyl cyanides of the formula VI,

$$\begin{array}{ccc}
 & & & VI \\
 & & & & VI \\
 & & & & & VI
\end{array}$$

are reacted with 3-amino-1,2,4-triazole of the formula III according to claim 6.

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8. (Currently amended) A fungicidal composition comprising a solid or liquid carrier and a compound of the formula I according to any of claims 1 to 5 claim 1.

- 9. (Currently amended) Seed comprising a compound of the formula I according to any of elaims 1 to 5 claim 1 in an amount of 1 to 1000 g per 100 kg.
- 10. (Currently amended) A method for controlling phytopathogenic harmful fungi wherein the fungi or the materials, plants, the soil or seed to be protected against fungal attack are treated with an effective amount of a compound of the formula I according to any of claims 1 to 5 claim 1.